10/581,545 - R1 - Brown et al. - SRNT - CAPLUS structure search

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NEWS 3	FEB 02	Simultaneous left and right truncation (SLART) added for CERAB, COMPUAB, ELCOM, and SOLIDSTATE
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NEWS 5	FEB 06	Patent sequence location (PSL) data added to USGENE
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NEWS 12	FEB 23	TOXCENTER updates mirror those of MEDLINE - more precise author group fields and 2009 MeSH terms
NEWS 13	FEB 23	Three million new patent records blast AEROSPACE into STN patent clusters
NEWS 14	FEB 25	USGENE enhanced with patent family and legal status display data from INPADOCDB
NEWS 15	MAR 06	INPADOCDB and INPAFAMDB enhanced with new display formats
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NEWS 17	MAR 11	ESBIOBASE reloaded and enhanced
NEWS 18	MAR 20	CAS databases on STN enhanced with new super role for nanomaterial substances
NEWS 19	MAR 23	CA/CAplus enhanced with more than 250,000 patent equivalents from China
NEWS 20	MAR 30	IMSPATENTS reloaded and enhanced
NEWS 21	APR 03	CAS coverage of exemplified prophetic substances enhanced
NEWS 22	APR 07	STN is raising the limits on saved answers
NEWS 23	APR 24	CA/CAplus now has more comprehensive patent assignee information
NEWS 24	APR 26	USPATFULL and USPAT2 enhanced with patent assignment/reassignment information
NEWS 25	APR 28	CAS patent authority coverage expanded
NEWS 26	APR 28	ENCOMPLIT/ENCOMPLIT2 search fields enhanced
NEWS 27	APR 28	Limits doubled for structure searching in CAS REGISTRY
NEWS 28	MAY 08	STN Express, Version 8.4, now available

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 NEWS 29  MAY 11  STN on the Web enhanced
 NEWS 30  MAY 11  BEILSTEIN substance information now available on
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 NEWS 31  MAY 14  DGENE, PCTGEN and USGENE enhanced with increased
                  limits for exact sequence match searches and
                  introduction of free HIT display format
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                  status data

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              AND CURRENT DISCOVER FILE IS DATED 06 APRIL 2009.

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=>
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Uploading A:\10.581545.R1.Brown et al..SRNT.CAPLUS

L1 STRUCTURE UPLOADED

=> d l1
L1 HAS NO ANSWERS
L1 STR

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

Structure attributes must be viewed using STN Express query preparation.

=> s l1 sss sam
SAMPLE SEARCH INITIATED 15:32:42 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 8 TO ITERATE

100.0% PROCESSED 8 ITERATIONS 0 ANSWERS
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 8 TO 329
PROJECTED ANSWERS: 0 TO 0

L2 0 SEA SSS SAM L1

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FULL SCREEN SEARCH COMPLETED - 176 TO ITERATE

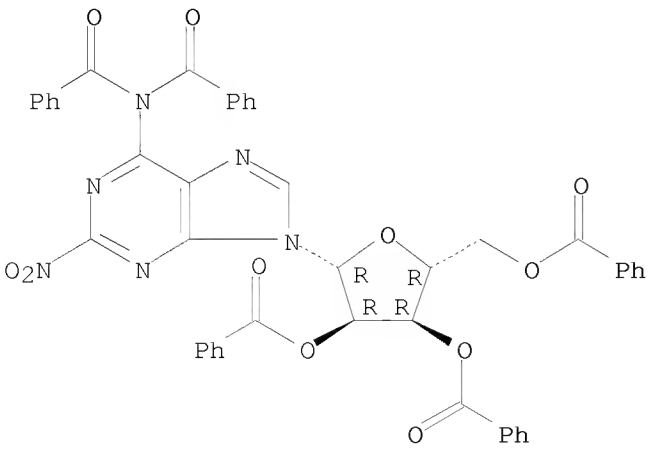
100.0% PROCESSED 176 ITERATIONS 6 ANSWERS
SEARCH TIME: 00.00.01

L3 6 SEA SSS FUL L1

=> d scan

L3 6 ANSWERS REGISTRY COPYRIGHT 2009 ACS on STN
IN Adenosine, N,N-dibenzoyl-2-nitro-, 2',3',5'-tribenzoate
MF C45 H32 N6 O11

Absolute stereochemistry.



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FULL ESTIMATED COST | 186.84 | 187.06

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FILE COVERS 1907 - 24 May 2009 VOL 150 ISS 22
FILE LAST UPDATED: 22 May 2009 (20090522/ED)
REVISED CLASS FIELDS (/NCL) LAST RELOADED: Feb 2009
USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Feb 2009

CAplus now includes complete International Patent Classification (IPC) reclassification data for the third quarter of 2008.

CAS Information Use Policies apply and are available at:

http://www.cas.org/legal/infopolicy.html

This file contains CAS Registry Numbers for easy and accurate

=> d his

(FILE 'HOME' ENTERED AT 15:30:57 ON 24 MAY 2009)

FILE 'REGISTRY' ENTERED AT 15:31:12 ON 24 MAY 2009
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L2 0 S L1 SSS SAM
L3 6 S L1 SSS FULL

FILE 'CAPLUS' ENTERED AT 15:33:15 ON 24 MAY 2009

=> s l3
L4 17 L3

=> d l3 ibib abs hitstr 1-17
YOU HAVE REQUESTED DATA FROM FILE 'REGISTRY' - CONTINUE? (Y)/N:nh
YOU HAVE REQUESTED DATA FROM FILE 'REGISTRY' - CONTINUE? (Y)/N:end

=> d l4 ibib abs hitstr 1-17

L4 ANSWER 1 OF 17 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2008:11519 CAPLUS
DOCUMENT NUMBER: 148:100840
TITLE: Preparation of nucleosides as therapeutic compounds for the treatment of pain and inflammation
INVENTOR(S): Higginbottom, Michael; Savory, Edward Daniel; Brown, Giles Albert; Horgan, Viet-Anh Anne; Chapman, Emma Jane
PATENT ASSIGNEE(S): Biovitrum AB, Swed.
SOURCE: PCT Int. Appl., 70pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2008000745	A2	20080103	WO 2007-EP56378	20070626
WO 2008000745	A3	20080417		
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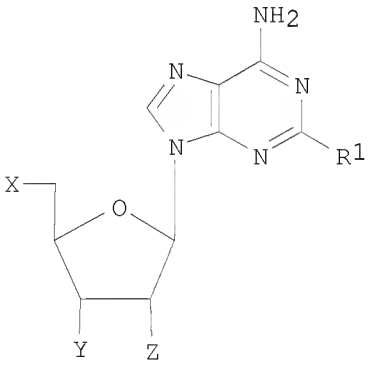
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA			
AU 2007263728	A1	20080103	AU 2007-263728	20070626
CA 2654375	A1	20080103	CA 2007-2654375	20070626
US 20080076776	A1	20080327	US 2007-823377	20070626
IN 2008KN04769	A	20090313	IN 2008-KN4769	20081125
PRIORITY APPLN. INFO.:			SE 2006-1398 A	20060627
			US 2006-837146P P	20060811
			WO 2007-EP56378 W	20070626

OTHER SOURCE(S): MARPAT 148:100840
GI



I

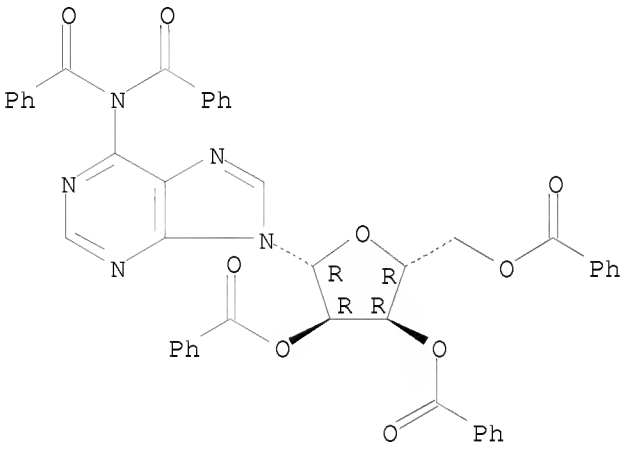
AB Nucleosides I, when X = Y = Z = OH, R1 is OCH2CF2CF3, phenoxy (substituted with 3-(4- trifluoromethylphenyl), 3,4-dichloro,

(3-trifluoromethyl,4-fluoro), (3-trifluoromethyl,4- chloro), (3-chloro, 4-cyano), or 3,5-bis(trifluoromethyl)), 1-piperazinyl(4-(3,4-dichlorophenyl)), Ph (substituted with 3,4-dichloro, 3,5-difluoro, 3,5-bis(trifluoromethyl) or 3,4,5-trifluoro) or 2-benzofuranyl; or when X = Y = OH and Z = OMe, R1 is OCH3, OCH2CHF2, OCH2cyclopentyl, O-(2,5-difluorophenyl) or (S)-sec-butylamino; or when X = H and Y = Z = OH, R1 is n-hexylamino or cyclopentylamino; or when (IV) X = Z = OH and Y = H, R1 is cyclopentylamino; were prepared for the treatment of pain and inflammation. Thus, nucleoside I (X = Y = Z = OH, R1 = CH2CF2CF3) was prepared and tested for the treatment of pain and inflammation.

IT 62374-23-6P 854158-99-9P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of nucleosides as therapeutic compds. for treatment of pain and inflammation)

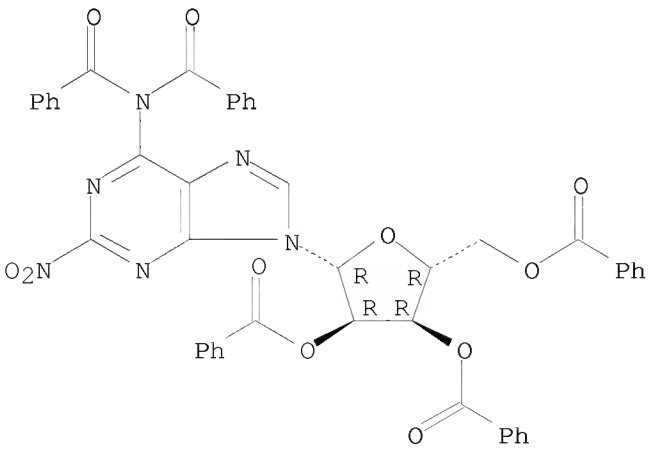
RN 62374-23-6 CAPLUS
CN Adenosine, N,N-dibenzoyl-, 2',3',5'-tribenzoate (CA INDEX NAME)

Absolute stereochemistry.



RN 854158-99-9 CAPLUS
CN Adenosine, N,N-dibenzoyl-2-nitro-, 2',3',5'-tribenzoate (CA INDEX NAME)

Absolute stereochemistry.

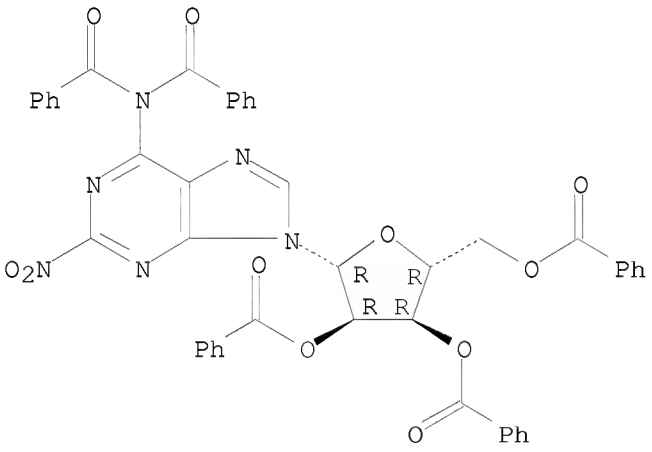


L4 ANSWER 2 OF 17 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2006:1143133 CAPLUS
DOCUMENT NUMBER: 146:82151
TITLE: A novel method for the introduction of fluorine into the purine 2-position: synthesis of 2-fluoroadenosine and a formal synthesis of the antileukemic drug fludarabine
AUTHOR(S): Braendvang, Morten; Gundersen, Lise-Lotte
CORPORATE SOURCE: Department of Chemistry, University of Oslo, Oslo, 0315, Norway
SOURCE: Synthesis (2006), (18), 2993-2995
CODEN: SYNTBF; ISSN: 0039-7881
PUBLISHER: Georg Thieme Verlag
DOCUMENT TYPE: Journal
LANGUAGE: English
OTHER SOURCE(S): CASREACT 146:82151

AB A novel method for the introduction of fluorine in the purine 2-position is described and employed in the synthesis of a potential antimycobacterial compound Also 2-fluoroadenosine has been synthesized for the first time from adenosine with perbenzoylated 2-nitroadenosine as a key intermediate. Mild reaction conditions are employed and few synthetic steps are required. The novel synthesis of fluoroadenosine can be regarded as a formal synthesis of the antileukemic drug fludarabine phosphate.

IT 854158-99-9P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(synthesis of 2-fluoroadenosine as a synthon toward a formal synthesis of the antileukemic drug fludarabine)
RN 854158-99-9 CAPLUS
CN Adenosine, N,N-dibenzoyl-2-nitro-, 2',3',5'-tribenzoate (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT: 23 THERE ARE 23 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 3 OF 17 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2005:1004549 CAPLUS
DOCUMENT NUMBER: 143:286636
TITLE: Preparation of nucleosides as adenosine receptors and used for the treatment of pain and inflammation

INVENTOR(S): Pritchard, Martyn; Ouzman, Jacqueline; Savory, Edward; Brown, Giles

PATENT ASSIGNEE(S): Cambridge Biotechnology Limited, UK

SOURCE: PCT Int. Appl., 81 pp.
CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005084653	A2	20050915	WO 2005-GB800	20050304
WO 2005084653	A3	20060518		
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RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
WO 2004079329	A2	20040916	WO 2004-GB902	20040305
WO 2004079329	A3	20041209		
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AU 2005218997	A1	20050915	AU 2005-218997	20050304
CA 2557285	A1	20050915	CA 2005-2557285	20050304
EP 1749016	A2	20070207	EP 2005-717878	20050304
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR				
CN 1946732	A	20070411	CN 2005-80007119	20050304
BR 2005008488	A	20070731	BR 2005-8488	20050304
JP 2007526291	T	20070913	JP 2007-501345	20050304
MX 2006010075	A	20070410	MX 2006-10075	20060904
NO 2006004365	A	20061122	NO 2006-4365	20060926
KR 2007004792	A	20070109	KR 2006-720304	20060929
IN 2006CN03674	A	20070706	IN 2006-CN3674	20061005
US 20080221060	A1	20080911	US 2007-598520	20071207

PRIORITY APPLN. INFO.:

GB 2004-5009	A	20040305
GB 2004-5012	A	20040305
WO 2004-GB902	A	20040305
GB 2004-12261	A	20040602
GB 2004-12262	A	20040602
GB 2004-13627	A	20040618
GB 2004-19718	A	20040906
GB 2004-20063	A	20040909
GB 2004-20615	A	20040916
GB 2003-5153	A	20030307
WO 2005-GB800	W	20050304

OTHER SOURCE(S): CASREACT 143:286636; MARPAT 143:286636

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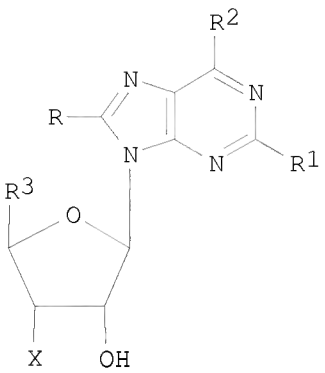


I

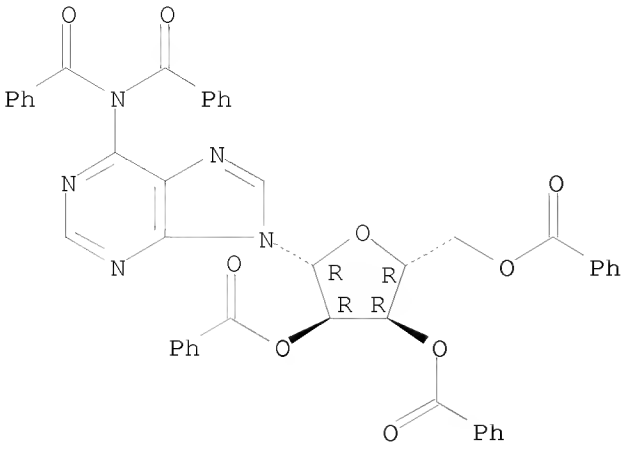
AB Nucleosides I, wherein X is H, OH; R is H, Me; R1 is H, alkoxy, OCH2-cyclopropyl, OCH2-cyclopentyl, phenoxy, OCH2CH2OH, OCH2CH2F2, (5-indanyl)oxy, alkylamino, cyclo-alkylamino, exo-norbornane, amino, phenylamino; R2 is NH2, CH2OH, NMe2, methylamino, isoamyl; R3 is CH2OH, amide, CH2NHCOPr-n, CH2NHCONHEt; were prepared and used for the treatment of pain and inflammation. Title nucleosides were prepared and used the treatment of pain associated with cancer, pancreatic pain, pain associated with HIV infection, chronic neuropathic pain, lower back pain, failed back surgery pain, back pain, post-operative pain, post phys. trauma pain, cardiac pain, chest pain, joint pain, neck pain, bowel pain, phantom limb pain, obstetric pain, acute herpes zoster pain, acute pancreatitis breakthrough pain, or for the prevention, treatment, or amelioration of neuropathic or other pain caused by, or associated with diabetic neuropathy, poly-neuropathy, fibromyalgia, myo-fascial pain syndrome, osteoarthritis, post herpetic neuralgia, rheumatoid arthritis, sciatica/lumbar radiculopathy, spinal stenosis, trigeminal neuralgia, renal colic, dysmenorhoea/endometriosis. Thus, I (R = H, R1 = OMe, R2 = NH1, R3 = CH2OH) was prepared and tested for the treatment of pain and inflammation.

IT 62374-23-6P 854158-99-9P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of nucleosides as adenosine receptors and used for the treatment of pain and inflammation)

RN 62374-23-6 CAPLUS

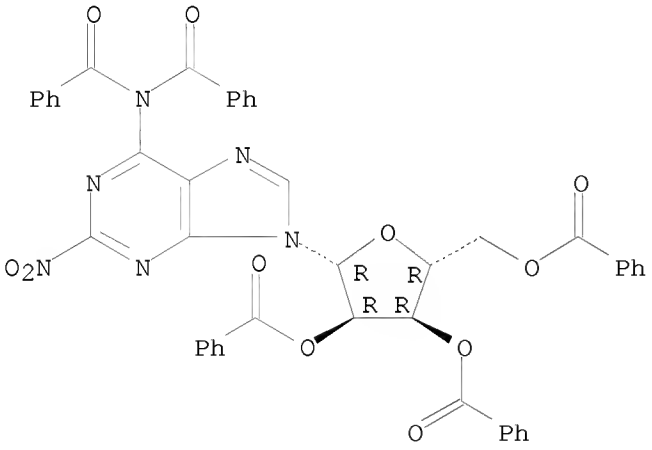
CN Adenosine, N,N-dibenzoyl-, 2',3',5'-tribenzoate (CA INDEX NAME)

Absolute stereochemistry.



RN 854158-99-9 CAPLUS
CN Adenosine, N,N-dibenzoyl-2-nitro-, 2',3',5'-tribenzoate (CA INDEX NAME)

Absolute stereochemistry.



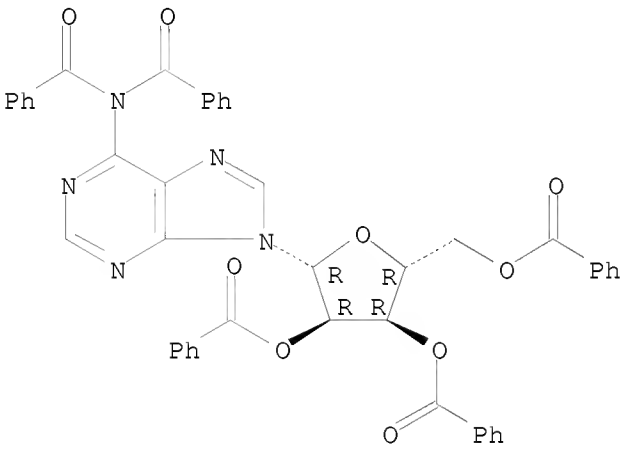
L4 ANSWER 4 OF 17 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2005:648167 CAPLUS
DOCUMENT NUMBER: 143:267187
TITLE: Nucleic Acid Related Compounds. 127. Selective N-Deacylation of N,O-Peracylated Nucleosides in Superheated Methanol
AUTHOR(S): Nowak, Ireneusz; Conda-Sheridan, Martin; Robins, Morris J.
CORPORATE SOURCE: Department of Chemistry and Biochemistry, Brigham Young University, Provo, UT, 84602-5700, USA
SOURCE: Journal of Organic Chemistry (2005), 70(18), 7455-7458
CODEN: JOCEAH; ISSN: 0022-3263
PUBLISHER: American Chemical Society
DOCUMENT TYPE: Journal
LANGUAGE: English
OTHER SOURCE(S): CASREACT 143:267187
AB Solns. of peracylated adenosine, cytidine, and related nucleoside derivs. undergo selective N-deacylation upon heating at elevated temps. (oil bath ≥ 105 °C) in methanol. An increase in the bulk of the

N-acyl group has little effect on the rate of N-deacylation but increases the N/O selectivity ratio. Extended heating is required for N-deacylation with aryl-carboxylic acid derivs. Contamination with acidic or basic reagent residues is avoided.

IT 62374-23-6P 863591-87-1P 863591-88-2P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(regioselective N-deacylation of per-acylated nucleosides in super-heated methanol)

RN 62374-23-6 CAPLUS
CN Adenosine, N,N-dibenzoyl-, 2',3',5'-tribenzoate (CA INDEX NAME)

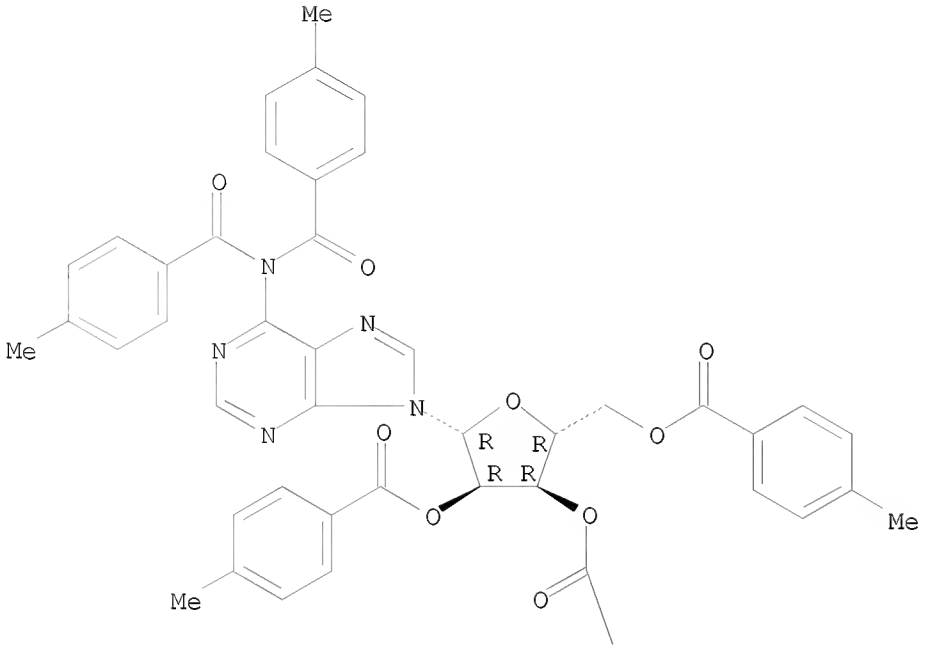
Absolute stereochemistry.



RN 863591-87-1 CAPLUS
CN Adenosine, N,N-bis(4-methylbenzoyl)-, 2',3',5'-tris(4-methylbenzoate) (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A



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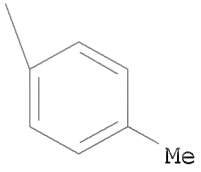
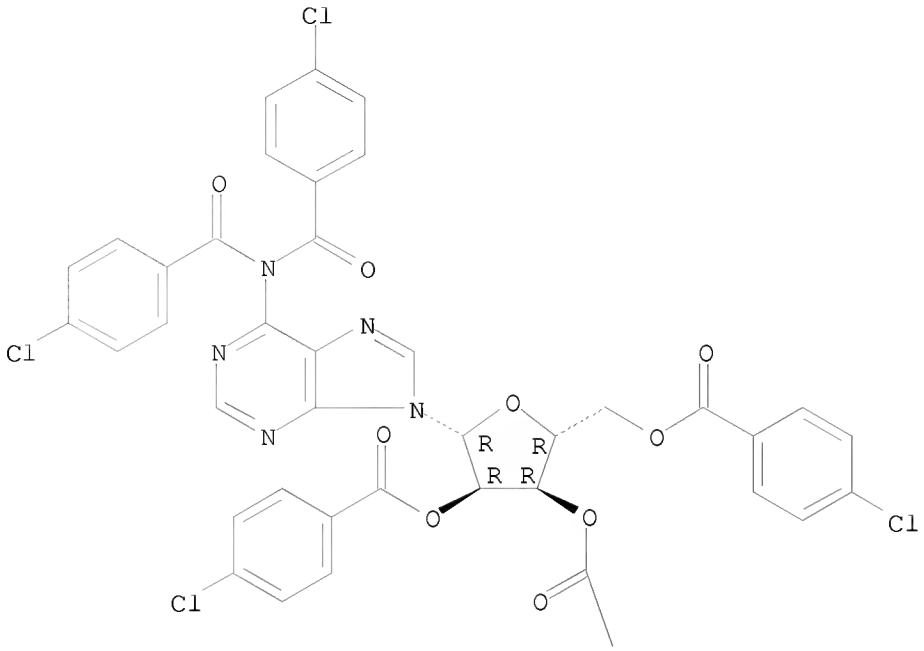


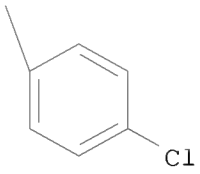
RN 863591-88-2 CAPLUS
CN Adenosine, N,N-bis(4-chlorobenzoyl)-, 2',3',5'-tris(4-chlorobenzoate) (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A



PAGE 2-A



REFERENCE COUNT: 24 THERE ARE 24 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 5 OF 17 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2005:540588 CAPLUS
DOCUMENT NUMBER: 143:60193
TITLE: Improved synthesis of 2-substituted adenosines via nitration reaction
INVENTOR(S): Brown, Giles Albert; Savory, Edward Daniel; Ouzman, Jacqueline Valerie Anne; Stoddart, Alison Margaret
PATENT ASSIGNEE(S): Cambridge Biotechnology Limited, UK
SOURCE: PCT Int. Appl., 32 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

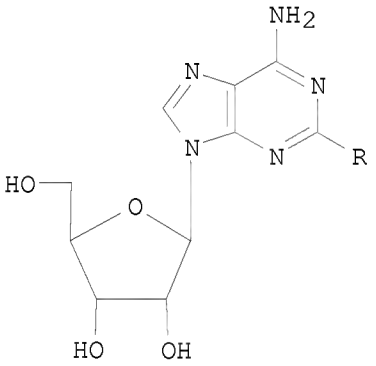
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005056571	A1	20050623	WO 2004-GB5090	20041203

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,

GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI,
NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY,
TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM,
AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK,
EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT,
RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML,
MR, NE, SN, TD, TG

AU 2004296242	A1	20050623	AU 2004-296242		20041203
CA 2552583	A1	20050623	CA 2004-2552583		20041203
EP 1694691	A1	20060830	EP 2004-805918		20041203
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK, IS					
CN 1886414	A	20061227	CN 2004-80035597		20041203
CN 100455591	C	20090128			
JP 2007513134	T	20070524	JP 2006-542018		20041203
NO 2006003109	A	20060905	NO 2006-3109		20060704
KR 2006125830	A	20061206	KR 2006-713388		20060704
IN 2006CN02462	A	20070608	IN 2006-CN2462		20060705
US 20090131651	A1	20090521	US 2008-581545		20081114
PRIORITY APPLN. INFO.:			GB 2003-28319	A	20031205
			GB 2003-28321	A	20031205
			WO 2004-GB5090	W	20041203

OTHER SOURCE(S): CASREACT 143:60193; MARPAT 143:60193
GI



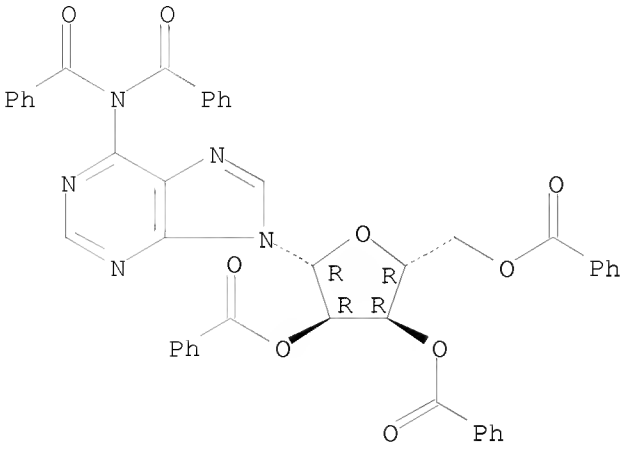
I

AB Synthesis of 2-substituted adenosines I, using 2-nitro penta-benzoyl adenosine, or 2-nitro penta-acetyl adenosine, as intermediate is described, wherein R = C1-6 alkoxy (straight or branched), a phenoxy group (unsubstituted, or mono-, or di-substituted by halo, amino, CF3, cyano, nitro, C1-6 alkyl, or C1-6 alkoxy), a benzyloxy group (unsubstituted, or mono-, or di-substituted by halo, amino, CF3, cyano, nitro, C1-6 alkyl, or C1-6 alkoxy), or a benzoyl group (unsubstituted, or mono-, or di-substituted by halo, amino, CF3-,cyano, nitro, C1-6 alkyl, or C1-6 alkoxy). The methods provide improved yield and purity of product. Thus, nucleoside spongosine was prepared from adenosine via benzoylation or acetylation followed by nitration and methoxylation reactions.

IT 62374-23-6P 854158-99-9P
RL: IMF (Industrial manufacture); RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(synthesis of 2-substituted adenosines via nitration reaction)

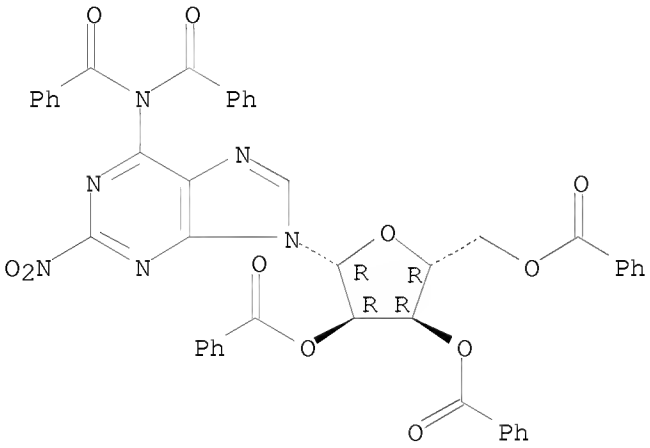
RN 62374-23-6 CAPLUS
CN Adenosine, N,N-dibenzoyl-, 2',3',5'-tribenzoate (CA INDEX NAME)

Absolute stereochemistry.



RN 854158-99-9 CAPLUS
CN Adenosine, N,N-dibenzoyl-2-nitro-, 2',3',5'-tribenzoate (CA INDEX NAME)

Absolute stereochemistry.

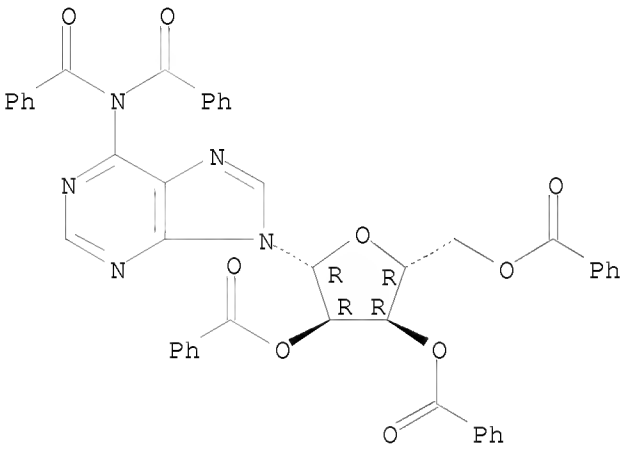


REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 6 OF 17 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 1996:94598 CAPLUS
DOCUMENT NUMBER: 124:261582
ORIGINAL REFERENCE NO.: 124:48483a
TITLE: Synthesis of N6,2',3',5'-tetrabenzoyl-β-D-adenosine catalyzed by metal iodides
AUTHOR(S): Nagai, Masashi; Matsutani, Takafumi; Mukaiyama, Teruaki
CORPORATE SOURCE: Faculty of Science, Science University of Tokyo, Tokyo, 162, Japan
SOURCE: Heterocycles (1996), 42(1), 57-63
CODEN: HTCYAM; ISSN: 0385-5414

PUBLISHER: Japan Institute of Heterocyclic Chemistry
DOCUMENT TYPE: Journal
LANGUAGE: English
OTHER SOURCE(S): CASREACT 124:261582
AB N-glycosidation of N6-benzoyl-N6,N9-bis(trimethylsilyl)adenine with Me 2,3,5-tri-O-benzoyl-β-D-ribofuranosyl carbonate was effectively promoted by several metal iodides and a desired coupling product, N6,2',3',5'-tetrabenzoyl-β-D-adenosine, was obtained in high yield when SbI3 or TeI4 was used as a catalyst.
IT 62374-23-6
RL: RCT (Reactant); RACT (Reactant or reagent)
(synthesis of tetrabenzoyladenosine catalyzed by metal iodides)
RN 62374-23-6 CAPLUS
CN Adenosine, N,N-dibenzoyl-, 2',3',5'-tribenzoate (CA INDEX NAME)

Absolute stereochemistry.



L4 ANSWER 7 OF 17 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 1990:210963 CAPLUS
DOCUMENT NUMBER: 112:210963
ORIGINAL REFERENCE NO.: 112:35457a,35460a
TITLE: Preparation of 2',5'-phosphorothioate oligoadenylates as virucides
INVENTOR(S): Suhadolnik, Robert J.; Pfleiderer, Wolfgang
PATENT ASSIGNEE(S): Temple University, USA
SOURCE: PCT Int. Appl., 74 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 8903683	A1	19890505	WO 1988-US3634	19881018
W: AU, BB, BG, BR, DK, FI, HU, JP, KP, KR, LK, MC, MG, MW, NO, RO, SD, SU				
RW: AT, BE, BJ, CF, CG, CH, CM, DE, FR, GA, GB, IT, LU, ML, MR, NL, SE, SN, TD, TG				
US 4924624	A	19900515	US 1987-112591	19871022
AU 8826280	A	19890523	AU 1988-26280	19881018
EP 389521	A1	19901003	EP 1988-909900	19881018

EP 389521	B1	19960320			
R: AT, BE, CH, DE, FR, GB, IT, LI, LU, NL, SE					
JP 04500795	T	19920213	JP 1988-509144		19881018
JP 2733777	B2	19980330			
EP 694559	A2	19960131	EP 1995-202077		19881018
EP 694559	A3	19960731			
EP 694559	B1	19990414			
R: AT, BE, CH, DE, FR, GB, IT, LI, LU, NL, SE					
AT 135579	T	19960415	AT 1988-909900		19881018
CA 1339953	C	19980714	CA 1988-580426		19881018
AT 178902	T	19990415	AT 1995-202077		19881018
US 5188897	A	19930223	US 1990-499109		19900326
US 5405939	A	19950411	US 1992-915771		19920716
US 5556840	A	19960917	US 1994-348419		19941202
PRIORITY APPLN. INFO.:			US 1987-112591	A	19871022
			EP 1988-909900	A3	19881018
			WO 1988-US3634	A	19881018
			US 1990-499118	B1	19900326
			US 1992-915771	A3	19920716

OTHER SOURCE(S): CASREACT 112:210963; MARPAT 112:210963
GI

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

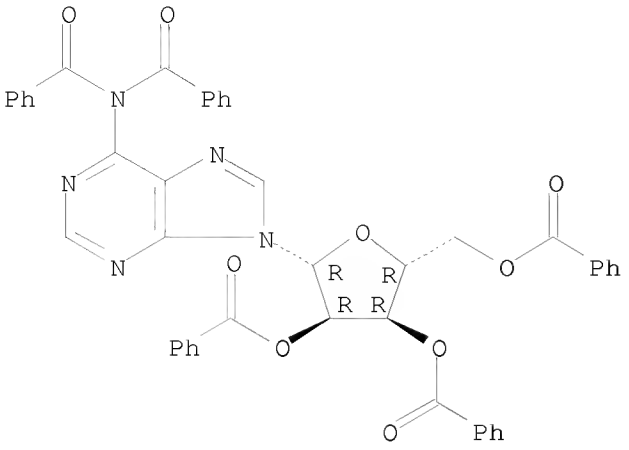
AB The optical isomers of the title compds. (I; m = 0-3; n = 1, 2) are prepared as medical and agrochem. virucides. 6-N-Benzoyl-3'-O-tert-butyldimethylsilyl-5'-O-monomethoxytrityl-(Sp)-P-thioadenylyl-2'-[O-(p-nitrophenylethyl)-5']-N-6-benzoyl-3'-O-tert-butyldimethylsilyl-(Sp)-P-thioadenylyl-2'-[O-(p-nitrophenylethyl)-5']-6-N-benzoyl-2',3'-bis-O-tert-butyldimethylsilyladenosine (preparation given) was detritylated with p-toluenesulfonic acid in CH2Cl2-MeOH (4:1), followed by TLC purification and reaction with 6-N-benzoyl-3'-O-tert-butyldimethylsilyl-5'-O-(4-ethoxytrityl)adenosine-2'-O-(p-nitrophenylethyl)octahydroazoninophosphoramidite (preparation given) in MeCN. 3-Nitro-1,2,4-triazole and S in pyridine were added to give the protected SpSpSp and RpSpSp tetramers, which were deblocked by stirring with DBU in pyridine, followed by neutralization and desilylation with Bu4NF in THF, to give (Rp)-P-thioadenyly-(2'→5')-(Sp)-P-thioadenylyl-(2'→5')-(Sp)-P-thioadenylyl-(2'→5')-adenosine (II) and its isomer. II (200 μM) inhibited the human immunodeficiency virus (HTLV-IIIBH-9) reverse transcriptase activity in vitro, as shown by a modification of the method of B. J. Poiesz, et al. (1980).

IT 62374-23-6P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation and debenzoylation of)

RN 62374-23-6 CAPLUS

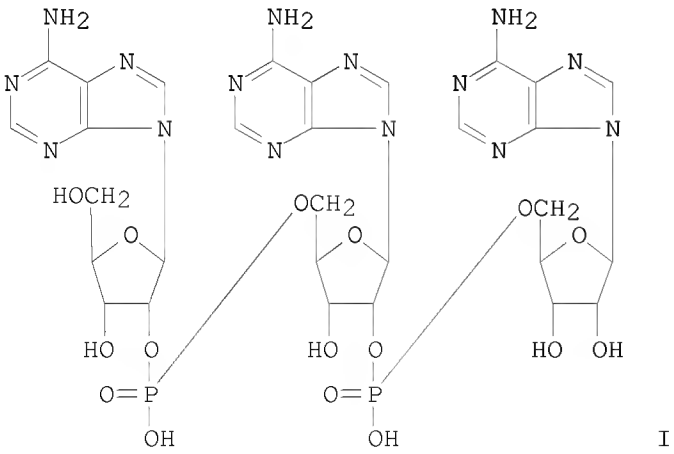
CN Adenosine, N,N-dibenzoyl-, 2',3',5'-tribenzoate (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 8 OF 17 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 1987:214280 CAPLUS
DOCUMENT NUMBER: 106:214280
ORIGINAL REFERENCE NO.: 106:34789a,34792a
TITLE: Solid-phase synthesis of 2'-5' oligoadenylate-2'-5' A core
AUTHOR(S): He, Binglin; Chen, Weizhu; Liu, Kellang; Zong, Jianchao; Li, Naihong
CORPORATE SOURCE: Dep. Chem., Nankai Univ., Tianjin, Peop. Rep. China
SOURCE: Scientia Sinica, Series B: Chemical, Biological, Agricultural, Medical & Earth Sciences (English Edition) (1986), 29(7), 686-97
CODEN: SSBSEF; ISSN: 0253-5823
DOCUMENT TYPE: Journal
LANGUAGE: English
GI



I

AB 2'-5' A core I was prepared on cross-linked polyacrylomorpholide resin as support by the phosphotriester approach using mesitylenesulfonyl chloride and N-methylimidazole as the condensing agents. Yields of the attachment

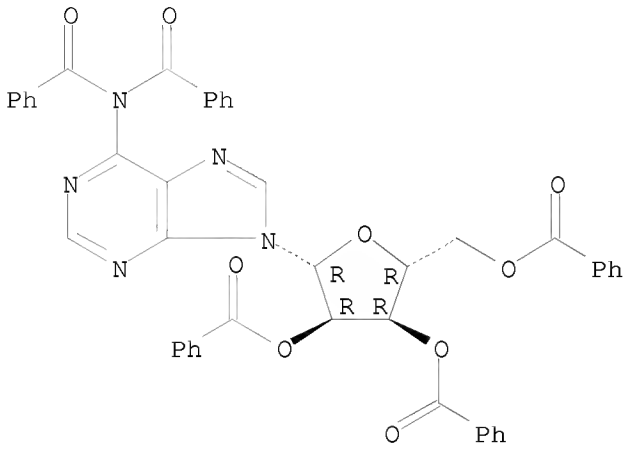
of the protected monomer to the support and subsequent 2 coupling reactions were 100, 60, and 91%, resp.

IT 62374-23-6P
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of, intermediate in solid-phase synthesis of 2'-5'-oligoadenylate)

RN 62374-23-6 CAPLUS

CN Adenosine, N,N-dibenzoyl-, 2',3',5'-tribenzoate (CA INDEX NAME)

Absolute stereochemistry.



L4 ANSWER 9 OF 17 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 1986:553466 CAPLUS

DOCUMENT NUMBER: 105:153466

ORIGINAL REFERENCE NO.: 105:24745a,24748a

TITLE: Regioselective protection of carbohydrate derivatives. Part 20. Simple, efficient 2'-O-deacylation of fully acylated purine and pyrimidine ribonucleosides through tert-butoxide

AUTHOR(S): Nishino, Shigeyoshi; Takamura, Hatsuko; Ishido, Yoshiharu

CORPORATE SOURCE: Fac. Sci., Tokyo Inst. Technol., Tokyo, 152, Japan

SOURCE: Tetrahedron (1986), 42(7), 1995-2004
CODEN: TETRAB; ISSN: 0040-4020

DOCUMENT TYPE: Journal

LANGUAGE: English

OTHER SOURCE(S): CASREACT 105:153466

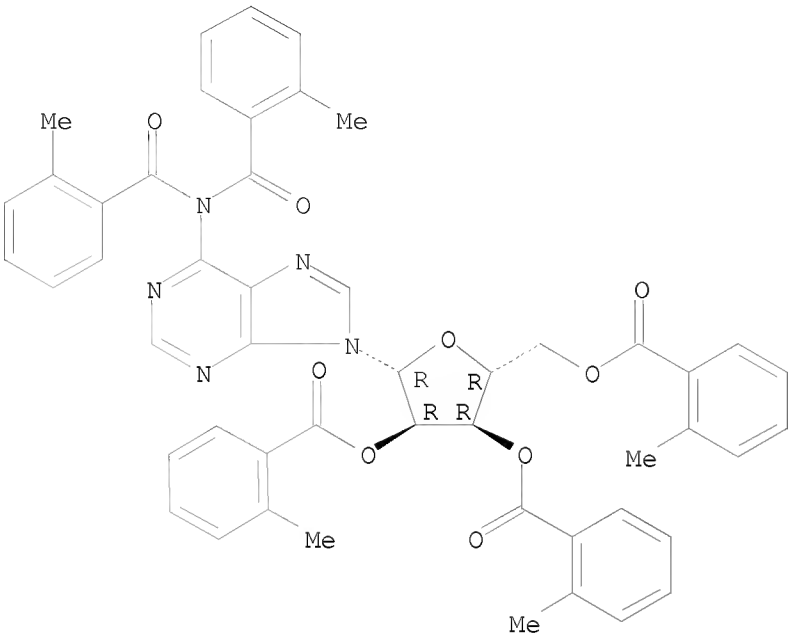
AB A simple treatment of fully aroylated purine and pyrimidine ribonucleosides with pulverized potassium tert-butoxide in THF or dichloromethane under a controlled condition gave a mixture of the corresponding di-O-aroyl derivs. in which 2'-OH derivs. are preponderant over 3'-OH derivs.; 3',5'-di-O-benzoyluridine, N4,3',5'-tribenzoylcytidine, N4,3',5'-tri-o-toluoylcytidine, N2,3',5'-tribenzoylguanosine, and N2-isobutyryl-3',5'-di-O-benzoylguanosine were obtained crystalline in 80%, 78%, 72%, 67%, and 65% yields, resp.

IT 104557-13-3P
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of, by regioselective O-dearoylation)

RN 104557-13-3 CAPLUS

CN Adenosine, N,N-bis(2-methylbenzoyl)-, 2',3',5'-tris(2-methylbenzoate) (9CI) (CA INDEX NAME)

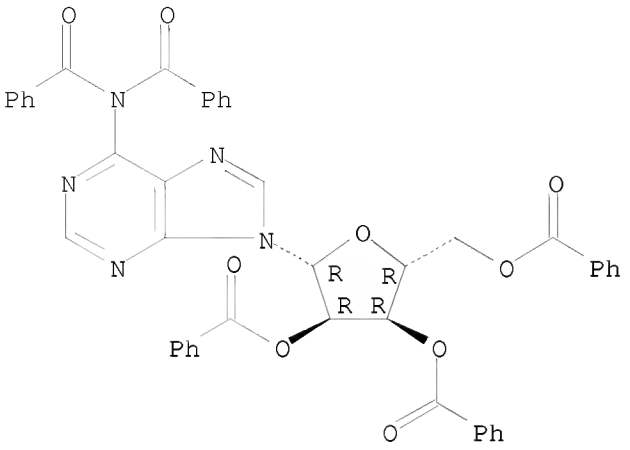
Absolute stereochemistry.



L4 ANSWER 10 OF 17 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 1985:62520 CAPLUS
DOCUMENT NUMBER: 102:62520
ORIGINAL REFERENCE NO.: 102:9825a,9828a
TITLE: Photobromination of carbohydrate derivatives. Part 8. Reaction of furanose derivatives with N-bromosuccinimide. X-ray molecular structure of 1-O-acetyl-2,5,6-tri-O-benzoyl-4-hydroxy-3,4-O-(α-succinimidobenzylidene)-β-D-galactofuranose
AUTHOR(S): Ferrier, Robert J.; Haines, Stephen R.; Gainsford, Graeme J.; Gabe, Eric J.
CORPORATE SOURCE: Dep. Chem., Victoria Univ. Wellington, Wellington, N. Z.
SOURCE: Journal of the Chemical Society, Perkin Transactions 1: Organic and Bio-Organic Chemistry (1972-1999) (1984), (8), 1683-7
CODEN: JCPRB4; ISSN: 0300-922X
DOCUMENT TYPE: Journal
LANGUAGE: English
GI For diagram(s), see printed CA Issue.
AB Photobromination of 1-O-acetyl-2,3,5,6-tetra-O-benzoyl-β-D-gluco- or -galactofuranose with NBS in refluxing CCl_4 for 0.9 h gave the title compound (I) in 74 and 64% yield, resp.; the structure of I was determined by x-ray crystallog. Similar reactions occurred with N-bromophthalimide and -acetamide. Reaction of 1-O-acetyl-2,3,5-tri-O-benzoyl-β-D-ribofuranose, pentabenzoyladenosine, and 1,2,3,4-tetra-O-acetyl-β-D-xylopyranose under these conditions gave the 4-, 4-, and 5-bromide, resp.
IT 62374-23-6
RL: RCT (Reactant); RACT (Reactant or reagent)
(bromination of, by NBS)
RN 62374-23-6 CAPLUS

CN Adenosine, N,N-dibenzoyl-, 2',3',5'-tribenzoate (CA INDEX NAME)

Absolute stereochemistry.

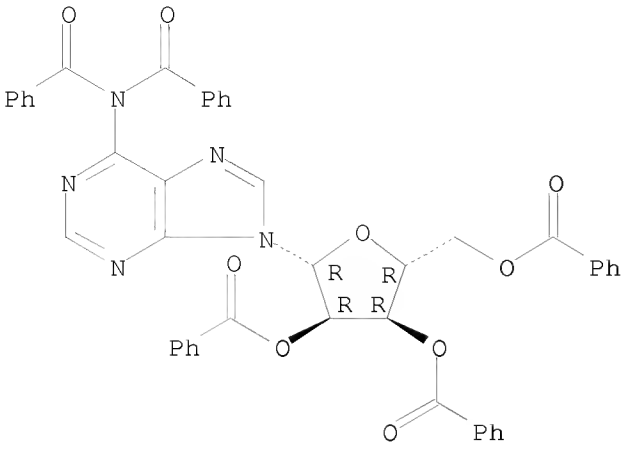


L4 ANSWER 11 OF 17 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 1981:620267 CAPLUS
DOCUMENT NUMBER: 95:220267
ORIGINAL REFERENCE NO.: 95:36765a,36768a
TITLE: Nucleosides. XXXVII. Synthesis and properties of 2'-O- and 3'-O-(tert-butyldimethylsilyl)-5'-O-(4-methoxytrityl)- and 2',3'-bis(O-tert-butyldimethylsilyl)ribonucleosides - starting materials for oligoribonucleotide syntheses
AUTHOR(S): Flockerzi, Dieter; Silber, Gunter; Charubala, Ramamurthy; Schlosser, Wilhelm; Varma, Rajendra Singh; Creegan, Frank; Pfleiderer, Wolfgang
CORPORATE SOURCE: Fak. Chem., Univ. Konstanz, Konstanz, D-7750, Fed. Rep. Ger.
SOURCE: Liebigs Annalen der Chemie (1981), (9), 1568-85
CODEN: LACHDL; ISSN: 0170-2041
DOCUMENT TYPE: Journal
LANGUAGE: German

AB The title nucleosides were prepared from adenosine, guanosine, cytidine, uridine, and 5,6-dihydrouridine. For example, N6-benzoyladenosine, prepared from adenosine in 2 steps, was treated with 4-methoxytrityl chloride and then silylated with Me3CSiMe2Cl to give 2'-O-(tert-butyldimethylsilyl)-, 3'-O-(tert-butyldimethylsilyl)-, and 2',3'-O-bis(tert-butyldimethylsilyl)-N6-benzoyl-5'-O-(4-methoxytrityl)adenosine (I). I was detritylated to give 2',3'-O-bis(tert-butyldimethylsilyl)-N6-benzoyladenosine. The compds. prepared were characterized by UV and 13C-NMR spectroscopy.

IT 62374-23-6P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation and partial debenzoylation of)
RN 62374-23-6 CAPLUS
CN Adenosine, N,N-dibenzoyl-, 2',3',5'-tribenzoate (CA INDEX NAME)

Absolute stereochemistry.

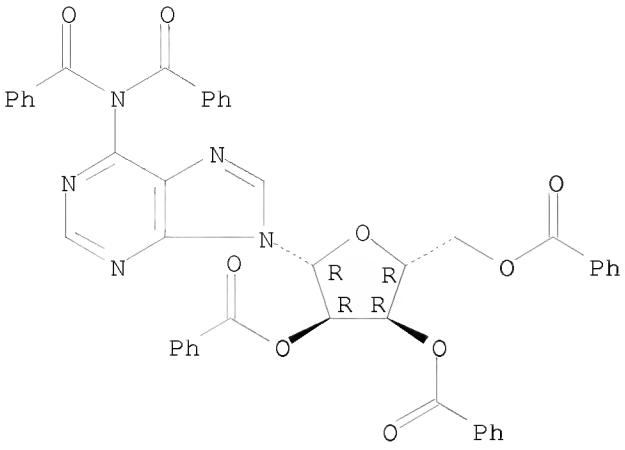


L4 ANSWER 12 OF 17 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 1980:59149 CAPLUS
DOCUMENT NUMBER: 92:59149
ORIGINAL REFERENCE NO.: 92:9815a,9818a
TITLE: Partial protection of carbohydrate derivatives. Part 3. Regioselective 2'-O-deacylation of fully acylated purine and pyrimidine ribonucleosides with hydrazine hydrate
AUTHOR(S): Ishido, Yoshiharu; Nakazaki, Nobuo; Sakairi, Nobuo
CORPORATE SOURCE: Dep. Chem., Tokyo Inst. Technol., Tokyo, Japan
SOURCE: Journal of the Chemical Society, Perkin Transactions 1: Organic and Bio-Organic Chemistry (1972-1999) (1979), (8), 2088-98
CODEN: JCPRB4; ISSN: 0300-922X
DOCUMENT TYPE: Journal
LANGUAGE: English

AB In 1:4 (volume/volume) AcOH-pyridine, partial O-deacylation of fully acylated purine and pyrimidine ribonucleosides upon hydrazinolysis was induced regioselectively in respect to 3 ester functions at the 2'-position to give the corresponding 2'-OH analogs in good yields. E.g., 3',5'-di-O-benzoyladenosine (70%), -inosine (52%), and -uridine (39%), N2-benzoyl-3',5'-diacetyl- (42%) and N2,3',5'-tribenzoylguanosine (63%) were isolated. 5'-O-Acylribonucleosides were prepared quant. using an excess of H2NNH2.H2O in 1:1 (volume/volume) CHCl3-MeOH and in pyridine. Hydrazinolysis of 3',5'-di-O-acetyl-2'-deoxyribonucleosides in pyridine gave both 5'- and 3'-O-acetyl-2'-deoxyribonucleosides (80-90% total yields). The 2'-O-acetyl group is more labile toward the nucleophile than the 3'-O-acetyl group. Possible factors involved in the regioselectivity of hydrazinolysis are discussed.

IT 62374-23-6
RL: RCT (Reactant); RACT (Reactant or reagent)
(partial deacylation of, with hydrazine)
RN 62374-23-6 CAPLUS
CN Adenosine, N,N-dibenzoyl-, 2',3',5'-tribenzoate (CA INDEX NAME)

Absolute stereochemistry.

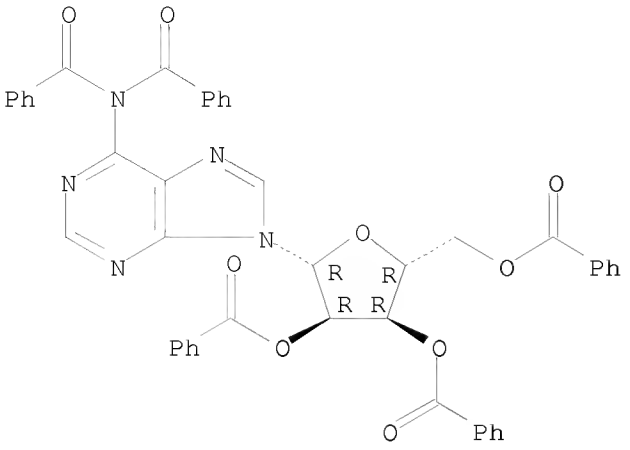


L4 ANSWER 13 OF 17 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 1979:575681 CAPLUS
DOCUMENT NUMBER: 91:175681
ORIGINAL REFERENCE NO.: 91:28351a,28354a
TITLE: Purinearabinosides
INVENTOR(S): Ishito, Ryoji; Sakairi, Nobuo
PATENT ASSIGNEE(S): Ajinomoto Co., Inc., Japan
SOURCE: Jpn. Kokai Tokkyo Koho, 3 pp.
CODEN: JKXXAF
DOCUMENT TYPE: Patent
LANGUAGE: Japanese
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 54073795	A	19790613	JP 1977-138208	19771117
PRIORITY APPLN. INFO.:			JP 1977-138208 A	19771117

AB Stirring a mixture of 1.58 g N6,N6,2',3',5'-pentabenzoyladenosine and 0.39 mL N2H4.H2O in 4:1 pyridine-AcOH 15 h at 70-5° gave 70% 3',5'-di-O-benzoyladenosine, which (1740 mg) was stirred with 10 mL Ac2O in Me2SO overnight at room temperature to give, after neutralization and extraction with AcOEt, a syrup, which was stirred with 150 mg NaBH4 in 1:1 C6H6-EtOH 2 h at 0°, filtered, concentrated, and stirred with a few drops of 2 N MeONa/MeOH 2 h at room temperature to give 537 mg 9-β-D-arabinofuranosyladenine. Similarly, 9-β-D-arabinofuranosylguanine was prepared from N2,2',3',5'-tetrabenzoylguanosine via N2,3',5'-tribenzoylguanosine.
IT 62374-23-6
RL: RCT (Reactant); RACT (Reactant or reagent)
(partial debenzoylation of)
RN 62374-23-6 CAPLUS
CN Adenosine, N,N-dibenzoyl-, 2',3',5'-tribenzoate (CA INDEX NAME)

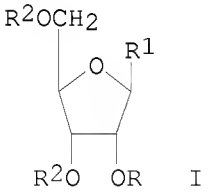
Absolute stereochemistry.



L4 ANSWER 14 OF 17 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 1978:615726 CAPLUS
DOCUMENT NUMBER: 89:215726
ORIGINAL REFERENCE NO.: 89:33541a,33544a
TITLE: Nucleoside derivatives
INVENTOR(S): Ishito, Ryoji; Nakasaki, Nobuo; Sakairi, Nobuo
PATENT ASSIGNEE(S): Ajinomoto Co., Inc., Japan
SOURCE: Jpn. Kokai Tokkyo Koho, 5 pp.
CODEN: JKXXAF
DOCUMENT TYPE: Patent
LANGUAGE: Japanese
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

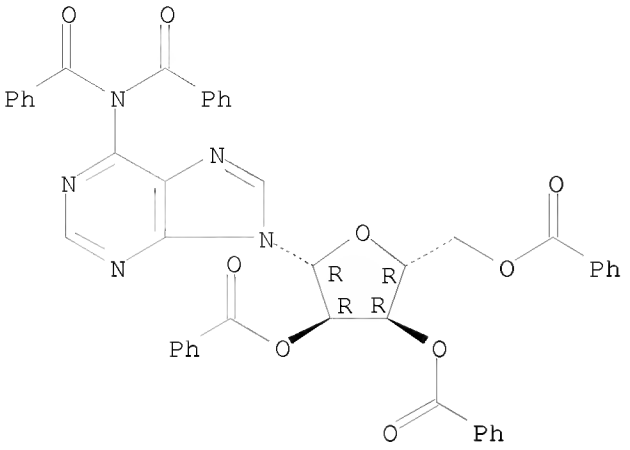
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 53084972	A	19780726	JP 1976-159283	19761229

PRIORITY APPLN. INFO.: JP 1976-159283 A 19761229
GI



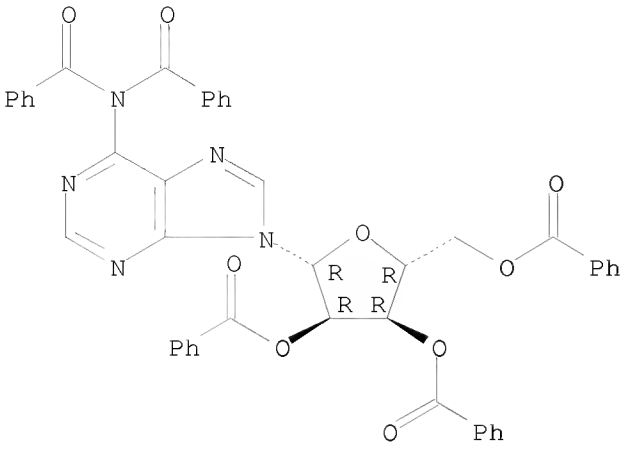
AB Nucleosides (I; R = H, R1 = nucleoside base, R2 = acyl) were prepared by reaction of I (R = R2 = acyl, R1 as before) with organic amines. Thus, a mixture of N6,N6,2',3',5'-pentabenzoyladenosine and N2H4.H2O in pyridine-AcOH was stirred 15 h at 70-5° to give 70% 3',5'-di-O-benzoyladenosine.
IT 62374-23-6
RL: RCT (Reactant); RACT (Reactant or reagent)
(partial debenzoylation of)
RN 62374-23-6 CAPLUS
CN Adenosine, N,N-dibenzoyl-, 2',3',5'-tribenzoate (CA INDEX NAME)

Absolute stereochemistry.



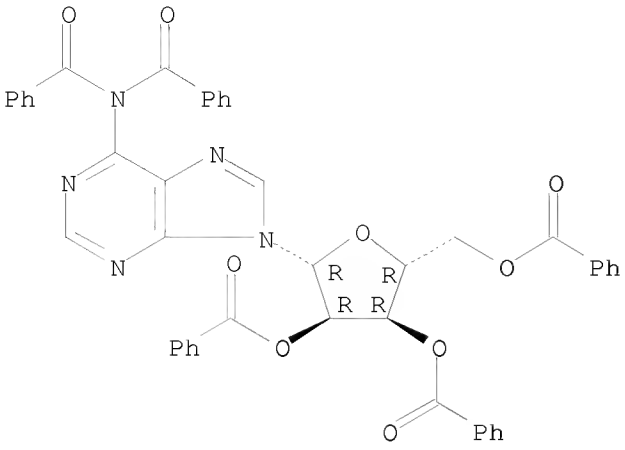
L4 ANSWER 15 OF 17 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 1977:468565 CAPLUS
DOCUMENT NUMBER: 87:68565
ORIGINAL REFERENCE NO.: 87:10941a,10944a
TITLE: Partial protection of carbohydrate derivatives. Part 1. Specific N-debenzoylation of fully benzoylated adenosine and cytidine with phenols and alcohols; active N-benzoyl groups
AUTHOR(S): Ishido, Yoshiharu; Nakazaki, Nobuo; Sakairi, Nobuo
CORPORATE SOURCE: Dep. Chem., Tokyo Inst. Technol., Tokyo, Japan
SOURCE: Journal of the Chemical Society, Perkin Transactions 1: Organic and Bio-Organic Chemistry (1972-1999) (1977), (6), 657-60
CODEN: JCPRB4; ISSN: 0300-922X
DOCUMENT TYPE: Journal
LANGUAGE: English
GI For diagram(s), see printed CA Issue.
AB Regiospecific N-debenzoylation of the adenosine derivs. I (R = H, Bz, R1 = Bz) (II and III, resp.) and the cytidine derivative IV (R = R1 = Bz) (V) occurred on treatment with various hydroxy compds. to give the corresponding nucleoside benzoates I (R = H, Bz, R1 = H) and IV (R = H, R1 = Bz), resp. The reaction is also a benzoylation of the hydroxy compds. by the nucleoside N-benzoyl groups; the order of activity is III > II > V. N-benzoylation of I (R = R1 = H) with 2,4-(O2N)2C6H3OBz and of IV (R = R1 = H) with III gave 83% II and 92% IV (R = Bz, R1 = H), resp.
IT 62374-23-6
RL: RCT (Reactant); RACT (Reactant or reagent)
(regiospecific debenzoylation of, by hydroxy compds.)
RN 62374-23-6 CAPLUS
CN Adenosine, N,N-dibenzoyl-, 2',3',5'-tribenzoate (CA INDEX NAME)

Absolute stereochemistry.



L4 ANSWER 16 OF 17 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 1977:140381 CAPLUS
DOCUMENT NUMBER: 86:140381
ORIGINAL REFERENCE NO.: 86:22061a,22064a
TITLE: A novel procedure for regioselective 2'-O-deacylation of fully acylated purine and pyrimidine ribonucleosides with hydrazine hydrate
AUTHOR(S): Ishido, Yoshiharu; Nakazaki, Nobuo; Sakairi, Nobuo
CORPORATE SOURCE: Fac. Sci., Tokyo Inst. Technol., Tokyo, Japan
SOURCE: Nucleic Acids Research, Special Publication (1976), 2(Symp. Nucleic Acids Chem., 4th, 1976), 25-8
CODEN: NARPD6; ISSN: 0309-1872
DOCUMENT TYPE: Journal
LANGUAGE: English
AB Hydrazinolyses of e.g. N6,N6,2',3',5'-pentabenzoyladenosine, N2,2',3',5'-tetrabenzoylguanosine, in AcOH-pyridine were regioselectively induced at 2' position of the alc. functions to give 63 and 44% of the corresponding 2'-hydroxyl derivs., resp. The procedure was also effective for partial debenzoylation of benzoylated uridine and cytidine although there was poorer regioselectivity. The treatment of N4,2',3',5'-tetraacetylcytidine was accompanied by no side reactions observed in that of the corresponding benzoate. The procedure was applied to the partial deacylation of 2'-deoxyribonucleoside acylates.
IT 62374-23-6
RL: RCT (Reactant); RACT (Reactant or reagent)
(deacylation of, hydrazine for)
RN 62374-23-6 CAPLUS
CN Adenosine, N,N-dibenzoyl-, 2',3',5'-tribenzoate (CA INDEX NAME)

Absolute stereochemistry.

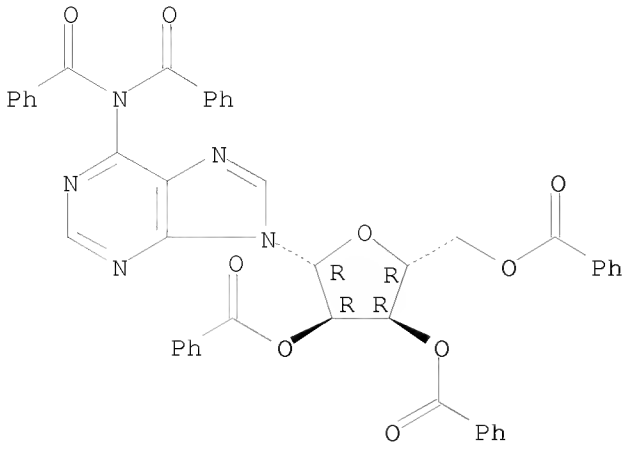


L4 ANSWER 17 OF 17 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 1977:106947 CAPLUS
DOCUMENT NUMBER: 86:106947
ORIGINAL REFERENCE NO.: 86:16885a,16888a
TITLE: Novel procedure for regioselective 2'-O-deacylation of fully acylated purine and pyrimidine ribonucleosides with hydrazine hydrate
AUTHOR(S): Ishido, Yoshiharu; Nakazaki, Nobuo; Sakairi, Nobuo
CORPORATE SOURCE: Dep. Chem., Tokyo Inst. Technol., Tokyo, Japan
SOURCE: Journal of the Chemical Society, Chemical Communications (1976), (20), 832-3
CODEN: JCCCAT; ISSN: 0022-4936
DOCUMENT TYPE: Journal
LANGUAGE: English

AB Hydrazinolysis of N6,N6,2',3',5'-pentabenzoyladenosine, N2,2',3',5'-tetrabenzoylguanosine, and 2',3',5'-tri-O-benzoylinosine with NH2NH2.H2O in AcOH-pyridine gave 68-70% of the corresponding 2'-OH derivs. Fully benzoylated uridine and cytidine were debenzoylated similarly but the regioselectivity observed was not as good. The same trend was observed in the hydrazinolysis of the corresponding acetates. E.g., hydrazinolysis of 2',3',5'-O-acetyladenosine gave 60% 3',5'-di-O-acetyladenosine.

IT 62374-23-6
RL: RCT (Reactant); RACT (Reactant or reagent)
(regioselective debenzoylation of, by hydrazinolysis)
RN 62374-23-6 CAPLUS
CN Adenosine, N,N-dibenzoyl-, 2',3',5'-tribenzoate (CA INDEX NAME)

Absolute stereochemistry.



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L3           6 S L1 SSS FULL

     FILE 'CAPLUS' ENTERED AT 15:33:15 ON 24 MAY 2009
L4          17 S L3

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     (FILE 'HOME' ENTERED AT 15:30:57 ON 24 MAY 2009)

     FILE 'REGISTRY' ENTERED AT 15:31:12 ON 24 MAY 2009
L1             STRUCTURE UPLOADED
L2           0 S L1 SSS SAM
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     FILE 'CAPLUS' ENTERED AT 15:33:15 ON 24 MAY 2009
L4          17 S L3

=> d his

     (FILE 'HOME' ENTERED AT 15:30:57 ON 24 MAY 2009)

     FILE 'REGISTRY' ENTERED AT 15:31:12 ON 24 MAY 2009
L1             STRUCTURE UPLOADED
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L4          17 S L3

=> d his

     (FILE 'HOME' ENTERED AT 15:30:57 ON 24 MAY 2009)
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     FILE 'REGISTRY' ENTERED AT 15:31:12 ON 24 MAY 2009
L1              STRUCTURE UPLOADED
L2            0 S L1 SSS SAM
L3            6 S L1 SSS FULL

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L4           17 S L3
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